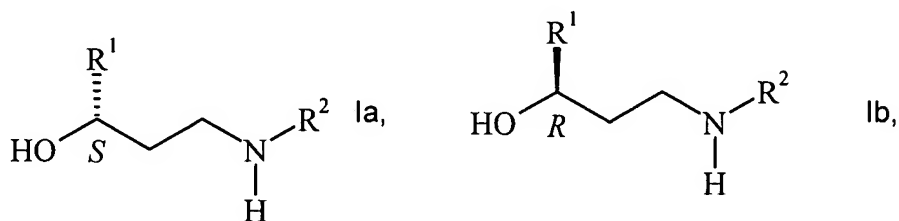


### Amendments To The Claims

This Listing Of Claims will replace all prior versions, and listings, of claims in the application:

#### Listing of Claims:

Claim 1 (Currently Amended): A process for the preparation of a salt of a carboxylic acid with an aminoalcohol of formula:



and/or

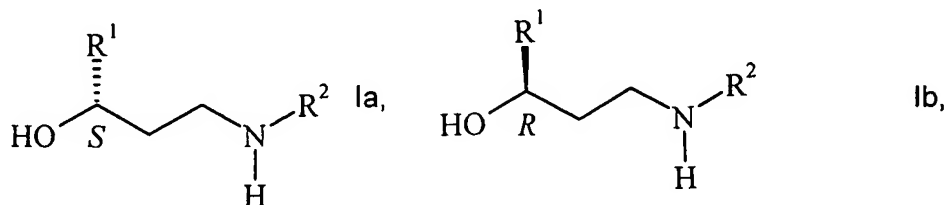
wherein  $R^1$  is selected from the group consisting of 2-thienyl, 2-furanyl, phenyl, 2-thienyl substituted with at least one halogen and/or at least one  $C_{1-4}$ -alkyl or  $C_{1-4}$ -alkoxy, 2-furanyl substituted with at least one halogen and/or at least one  $C_{1-4}$ -alkyl or  $C_{1-4}$ -alkoxy, and phenyl substituted with at least one halogen and/or at least one  $C_{1-4}$ -alkyl or  $C_{1-4}$ -alkoxy, and wherein  $R^2$  is selected from the group consisting of  $C_{1-4}$ -alkyl, phenyl,  $C_{1-4}$ -alkyl substituted with at least one halogen and/or at least one  $C_{1-4}$ -alkyl or  $C_{1-4}$ -alkoxy, and phenyl substituted with at least one halogen and/or at least one  $C_{1-4}$ -alkyl or  $C_{1-4}$ -alkoxy, comprising asymmetrically hydrogenating a salt of a carboxylic acid with an aminoketone of formula:



wherein  $R^1$  and  $R^2$  are as defined above,

in the presence of a catalytic amount of a catalyst comprising a transition metal complex of a diphosphine ligand.

Claim 2 (Currently Amended): A ~~The process of Claim 1,~~ comprising preparing for ~~the preparation of~~ a salt of a carboxylic acid with an aminoalcohol of formula:




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and/or

wherein R<sup>1</sup> is selected from the group consisting of 2-thienyl, 2-furanyl, phenyl, 2-thienyl substituted with at least one halogen and/or at least one C<sub>1-4</sub>-alkyl or C<sub>1-4</sub>-alkoxy, 2-furanyl substituted with at least one halogen and/or at least one C<sub>1-4</sub>-alkyl or C<sub>1-4</sub>-alkoxy, and phenyl substituted with at least one halogen and/or at least one C<sub>1-4</sub>-alkyl or C<sub>1-4</sub>-alkoxy, and wherein R<sup>2</sup> is C<sub>1-4</sub>-alkyl, phenyl, C<sub>1-4</sub>-alkyl substituted with at least one halogen and/or at least one C<sub>1-4</sub>-alkyl or C<sub>1-4</sub>-alkoxy, and phenyl substituted with at least one halogen and/or at least one C<sub>1-4</sub>-alkoxy, comprising by asymmetrically hydrogenating a salt of a carboxylic acid, wherein the carboxylic acid is selected from the group consisting of substituted C<sub>1-18</sub>-alkanoic acids, substituted monocyclic aromatics acids and substituted bicyclic acids, with an aminoketone of formula:



wherein R<sup>1</sup> and R<sup>2</sup> are as defined above,

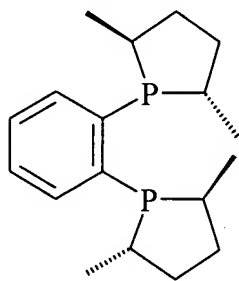
in the presence of a catalytic amount of a catalyst comprising a transition metal complex of a diphosphine ligand.

Claim 3 (Previously Presented): The process of claim 2, wherein R<sup>1</sup> is 2-thienyl, or 2-thienyl substituted with at least one halogen, and R<sup>2</sup> is methyl or ethyl.

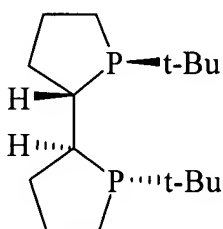
Claim 4 (Original): The process of claim 3, wherein the compound of formula II is selected from the group consisting of (S)-(-)-3-*N*-methylamino-1-(2-thienyl)-1-propanol, (S)-(-)-3-*N*-methyl-amino-1-(3-chloro-2-thienyl)-1-propanol, (R)-(+)-3-*N*-methylamino-1-(2-thienyl)-1-propanol and (R)-(+)-3-*N*-methylamino-1-(3-chloro-2-thienyl)-1-propanol.

Claim 5 (Currently Amended): The process of claim 4, wherein the transition metal is selected from the group consisting of rhodium, ruthenium and ~~or~~ iridium.

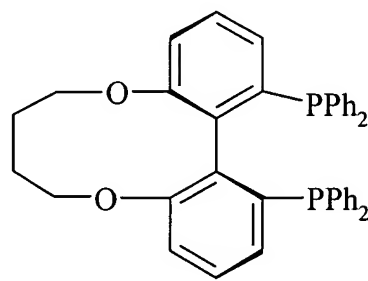
Claim 6 (Previously Presented): The process of claim 7, wherein the diphosphine ligand is selected from the group consisting of:



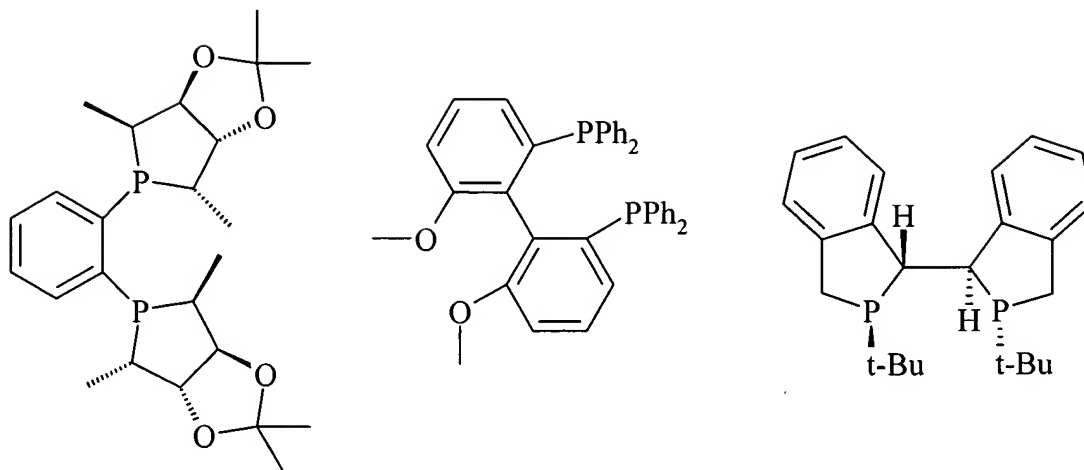
(S,S)-"Me-DuPhos",



(R,R,S,S)-"TangPhos",



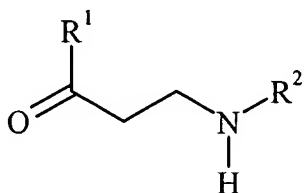
(S)-"C4-TunePhos",



(*S,S,S,S*)-"Me-KetalPhos", (*S*) and (*R*)-"MeO-BiPhep", and "(*R<sub>P</sub>*,*R<sub>P</sub>*,*S<sub>C</sub>*,*S<sub>C</sub>*)-DuanPhos".

Claim 7 (Previously Presented): The process of claim 6, wherein the compound of formulae Ia and/or Ib is obtained from its corresponding salt with a carboxylic acid by hydrolysis in the presence of an alkali metal hydroxide or an alkaline earth hydroxide.

Claim 8 (Withdrawn): A salt of a carboxylic acid with an aminoketone of the formula:



II,

wherein  $R^1$  is 2-thienyl or 2-furanyl, each optionally substituted with one or more halogen atoms and/or one or more  $C_{1-4}$ -alkyl or  $C_{1-4}$ -alkoxy groups, and wherein  $R^2$  is  $C_{1-4}$ -alkyl or phenyl, each optionally substituted with one or more halogen atoms and/or one or more  $C_{1-4}$ -alkyl or  $C_{1-4}$ -alkoxy groups.

Claim 9 (Withdrawn): The salt of claim 8, wherein the acid is selected from the group consisting of  $C_{1-18}$ -alkanoic acids,  
(-)-2,3:4,6-di-O-isopropylidene-2-keto-L-gulonic acid,

(+)-2,3:4,6-di-O-isopropylidene-2-keto-D-gulonic acid, 2-keto-L-gulonic acid, 2-keto-D-gulonic acid, L-aspartic acid, D-aspartic acid, DL-aspartic acid, benzoic acid, 3-methyl-benzoic acid, salicylic acid, 1-naphthalene carboxylic acid and 2-naphthalenecarboxylic acid.

Currently 10 (Withdrawn): A salt of a carboxylic acid with an aminoalkohol of the formula:



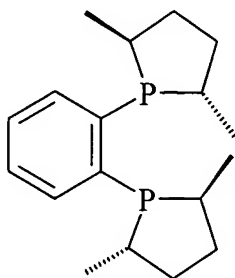
wherein  $\text{R}^1$  is 2-furanyl or phenyl, each optionally substituted with one or more halogen atoms and/or one or more  $\text{C}_{1-4}$ -alkyl or  $\text{C}_{1-4}$ -alkoxy groups, and wherein  $\text{R}^2$  is  $\text{C}_{1-4}$ -alkyl or phenyl, each optionally substituted with one or more halogen atoms and/or one or more  $\text{C}_{1-4}$ -alkyl or  $\text{C}_{1-4}$ -alkoxy groups, with the exception of salts, wherein the acid is (–)-2,3:4,6-di-O-isopropylidene-2-keto-L-gulonic acid or (+)-2,3:4,6-di-O-isopropylidene-2-keto-D-gulonic acid.

Claim 11 (Previously Presented): The process of claim 1, wherein the transitional metal complex of a diphosphine ligand is a transitional metal complex of an aryldiphosphine ligand or a biaryldiphosphine ligand.

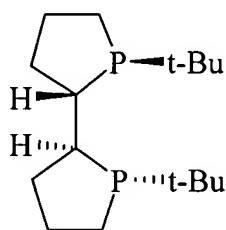
Claim 12 (Previously Presented): The process of claim 1, wherein  $\text{R}^1$  is 2-thienyl, optionally substituted with one or more halogen atoms, and  $\text{R}^2$  is methyl or ethyl.

Claim 13 (Previously Presented): The process of claim 1, wherein the transition metal is rhodium.

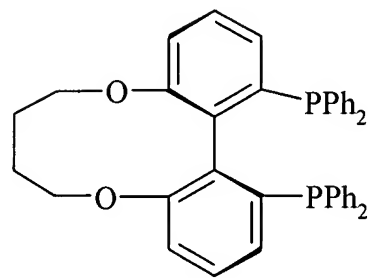
Claim 14 (Previously Presented): The process of claim 1, wherein the diphosphine ligand is selected from the group consisting of:



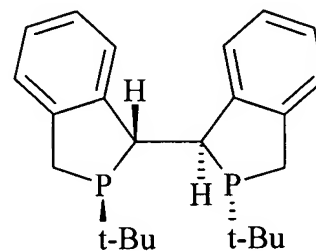
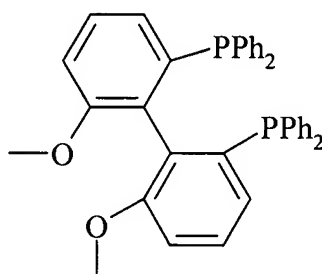
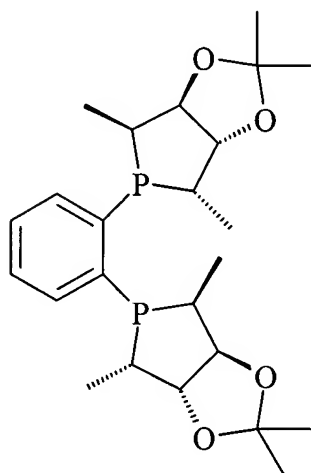
(*S,S*)-"Me-DuPhos",



(*R,R,S,S*)-"TangPhos",

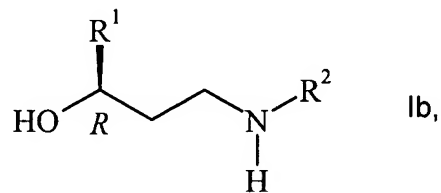
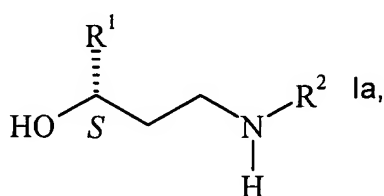


(*S*)-"C4-TunePhos",



(*S,S,S,S*)-"Me-KetalPhos", (*S*) and (*R*)-"MeO-BiPhep", and "(*R<sub>P</sub>*,*R<sub>P</sub>*,*S<sub>C</sub>*,*S<sub>C</sub>*)-DuanPhos".

Claim 15 (Currently Amended): A ~~The process of claim 1, wherein for the~~  
preparation of a salt of a carboxylic acid with an aminoalcohol of formula:

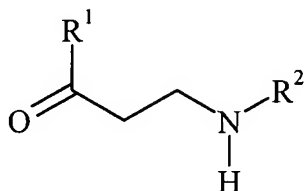


and/or

wherein R<sup>1</sup> is selected from the group consisting of 2-thienyl, 2-furanyl, phenyl,

2-thienyl substituted with at least one halogen and/or at least one C<sub>1-4</sub>-alkyl or C<sub>1-4</sub>-alkoxy, 2-furanyl substituted with at least one halogen and/or at least one C<sub>1-4</sub>-alkyl or C<sub>1-4</sub>-alkoxy, and phenyl substituted with at least one halogen and/or at least one C<sub>1-4</sub>-alkyl or C<sub>1-4</sub>-alkoxy, and wherein R<sup>2</sup> is selected from the group consisting of C<sub>1-4</sub>-alkyl, phenyl, C<sub>1-4</sub>-alkyl substituted with at least one halogen and/or at least one C<sub>1-4</sub>-alkyl or C<sub>1-4</sub>-alkoxy, and phenyl substituted with at least one halogen and/or at least one C<sub>1-4</sub>-alkyl or C<sub>1-4</sub>-alkoxy,  
comprising:

(i) asymmetrically hydrogenating a salt of a carboxylic acid with an aminoketone of formula:



II,

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wherein R<sup>1</sup> and R<sup>2</sup> are as defined above,

in the presence of a catalytic amount of a catalyst comprising a transition metal complex of a diphosphine ligand; and

(ii) obtaining a compound ~~the compounds of formulae Ia and/or Ib is obtained from its~~ corresponding salt with a carboxylic acid by hydrolysis of said corresponding salt in the presence of an alkali metal hydroxide or an alkaline metal hydroxide.

Claim 16 (Previously Presented): The process of claim 2, wherein the substituted C<sub>1-18</sub>-alkanoic acid is substituted with at least one C<sub>1-6</sub>-alkyl, C<sub>1-6</sub>-alkoxy, aryl, amino, protected carbonyl, halogen, hydroxyl or further carboxylic.

Claim 17 (Currently Amended): The process of claim 2, wherein the substituted

monocyclic aromatic acid is substituted with at least one member selected from the group consisting of C<sub>1-6</sub>-alkyl, C<sub>1-6</sub>alkoxy, halogen and ~~or~~ hydroxyl.

Claim 18 (Previously Presented): The process of claim 2, wherein the substituted bicyclic aromatic acid is substituted with at least one member selected from the group consisting of C<sub>1-6</sub>alkyl, C<sub>1-6</sub>-alkoxy, halogen and hydroxyl.

Claim 19 (Previously Presented): The process of ~~Claim~~ claim 1; wherein the catalyst is present in a catalytic amount.

Claim 20 (Previously Presented): The process of ~~Claim~~ claim 1, wherein the carboxylic acid is a monocarboxylic acid.